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- (ii) *Indications for use.* Treatment of canine urinary tract infections associated with *Escherichia coli* and *Proteus mirabilis.*
- (iii) *Limitations.* For subcutaneous use only. Treatment should be repeated at 24-hour intervals, continued for 48 hours after clinical signs have disappeared, for 5 to 14 days. Do not use in animals found to be hypersensitive to the drug. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (7) Sheep—(i) Amount. 0.5 to 1.0 milligram per pound (1.1 to 2.2 milligrams per kilogram) of body weight.
- (ii) Indications for use. For treatment of sheep respiratory disease (pneumonia) associated with Pasteurella haemolytica and/or P. multocida.
- (iii) *Limitations*. For intramuscular use only. Treatment should be repeated at 24 hour intervals for a total of 3 consecutive days. Additional treatments may be given on days 4 and 5 for animals which do not show satisfactory response. Use of dosages in excess of those indicated or by unapproved routes of administration may result in illegal residues in tissues. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (8) Goats—(i) Amount. 0.5 to 1.0 milligram per pound of body weight by intramuscular injection at 24-hour intervals for a total of 3 consecutive days. Additional treatments may be given on days 4 and 5 for animals that do not show satisfactory response.
- (ii) *Indications for use.* For treatment of caprine respiratory disease (goat pneumonia) associated with *Pasteurella haemolytica* and *P. multocida*.
- (iii) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[53 5369, Feb. 24, 1988, as amended at 55 FR 13768, Apr. 12, 1990; 56 FR 12119, Mar. 22, 1991; 57 FR 41862, Sept. 14, 1992; 59 FR 41666, Aug. 15, 1994; 59 FR 54518, Nov. 1, 1994; 60 FR 51719, Oct. 3, 1995; 61 FR 35130, July 5, 1996; 61 FR 66583, Dec. 18, 1996; 66 FR 21283, Apr. 30, 2001; 66 FR 32540, June 15, 2001; 69 FR 47362, Aug. 5, 2004]

§ 522.314 Ceftiofur hydrochloride.

(a) Specifications. Each milliliter of suspension contains ceftiofur hydro-

- chloride equivalent to 50 milligrams (mg) of ceftiofur.
- (b) Sponsor. See No. 000009 in \$510.600(c) of this chapter.
- (c) Related tolerances. See §556.113 of this chapter.
- (d) Conditions of use. (1) Swine—(i) Amount. 3 to 5 mg per kilogram (/kg) of body weight by intramuscular injection. Treatment should be repeated at 24-hour intervals for a total of 3 consecutive days.
- (ii) Indications for use. For treatment and control of swine bacterial respiratory disease (swine bacterial pneumonia) associated with Actinobacillus (Haemophilus) pleuropneumoniae, Pastureurella multocida, Salmonella choleraesuis, and Streptococcus suis Type
- (iii) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian. Treated swine must not be slaughtered for 4 days following the last treatment.
- (2) Cattle—(i) Dosage. 1.1 to 2.2 mg/kg of body weight by intramuscular or subcutaneous injection, at 24-hour intervals for 3 to 5 consecutive days. For bovine respiratory disease, 2.2 mg/kg of body weight may be administered twice at a 48-hour interval. For acute metritis, administer 2.2 mg/kg of body weight daily for 5 consecutive days.
- (ii) Indications for use. For treatment of bovine respiratory disease (BRD, shipping fever, pneumonia) associated with Mannheimia spp. (Pasteurella multocida, haemolytica). Pand Haemophilus somnus; acute bovine interdigital necrobacillosis (foot rot. pododermatitis) associated with Fusobacterium necrophorum Bacteroides melaninogenicus; and acute metritis (0 to 14 days post partum) associated with bacteria susceptible to ceftiofur.
- (iii) *Limitations.* Do not slaughter treated cattle for 48 hours (2 days) after last treatment. A withdrawal period has not been established in preruminating calves. Do not use in calves to be processed for veal. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[61 FR 29479, June 11, 1996, as amended at 63 FR 53578, Oct. 6, 1998; 67 FR 45901, July 11, 2002; 69 FR 47362, Aug. 5, 2004]